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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/936,565	02/04/2002	John J. Sauk	UNIMD 4	7145
23599 7590 10/19/2006 MILLEN, WHITE, ZELANO & BRANIGAN, P.C. 2200 CLARENDON BLVD.			EXAMINER	
			YAEN, CHRIS	YAEN, CHRISTOPHER H
SUITE 1400	DON BEVD.		ART UNIT	PAPER NUMBER
ARLINGTON,	VA 22201		1643	

DATE MAILED: 10/19/2006

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant/s)	
	Application No.	Applicant(s)	
Office Action Summary	09/936,565	SAUK, JOHN J.	
Office Action Summary	Examiner	Art Unit	
	Christopher H. Yaen	1643	
The MAILING DATE of this communication ap Period for Reply	pears on the cover sheet with	the correspondence address	
A SHORTENED STATUTORY PERIOD FOR REPL WHICHEVER IS LONGER, FROM THE MAILING D.  - Extensions of time may be available under the provisions of 37 CFR 1. after SIX (6) MONTHS from the mailing date of this communication.  - If NO period for reply is specified above, the maximum statutory period Failure to reply within the set or extended period for reply will, by statut Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	DATE OF THIS COMMUNICA 136(a). In no event, however, may a reply will apply and will expire SIX (6) MONTH: e, cause the application to become ABAN	TION.  be timely filed  from the mailing date of this communication.  DONED (35 U.S.C. § 133).	
Status			
1) Responsive to communication(s) filed on 28 J	l <u>uly 2006</u> .		
2a) This action is <b>FINAL</b> . 2b) ☐ This	s action is non-final.		
3) Since this application is in condition for allowa	ance except for formal matters	, prosecution as to the merits is	
closed in accordance with the practice under	Ex parte Quayle, 1935 C.D. 1	1, 453 O.G. 213.	
Disposition of Claims			
<ul> <li>4)  Claim(s) 1-19,24-26 and 29-38 is/are pending 4a) Of the above claim(s) 1-19 is/are withdraw</li> <li>5)  Claim(s) is/are allowed.</li> <li>6)  Claim(s) 24-26 and 29-38 is/are rejected.</li> <li>7)  Claim(s) 34 is/are objected to.</li> <li>8)  Claim(s) are subject to restriction and/o</li> </ul>	n from consideration.		
Application Papers			
9) The specification is objected to by the Examine 10) The drawing(s) filed on is/are: a) accomplicant may not request that any objection to the Replacement drawing sheet(s) including the correct 11) The oath or declaration is objected to by the Examine 11.	cepted or b) objected to by drawing(s) be held in abeyance ction is required if the drawing(s)	. See 37 CFR 1.85(a). is objected to. See 37 CFR 1.121(d).	
Priority under 35 U.S.C. § 119			
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of:  1. Certified copies of the priority document 2. Certified copies of the priority document 3. Copies of the certified copies of the priority application from the International Burea * See the attached detailed Office action for a list	ts have been received. ts have been received in App prity documents have been re tu (PCT Rule 17.2(a)).	lication No ceived in this National Stage	
Attachment(s)	4) 🗔 Intention Surre	(DTO 442)	
1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date		lail Date mal Patent Application	

### **DETAILED ACTION**

Re: Sauk J

1. The amendment filed 7/28/2006 is acknowledged and entered into the record.

Accordingly, claims 20-23 and 27-28 are canceled without prejudice or disclaimer, and

claims 33-38 are newly added.

2. Claims 1-19,24-26,29-38 are pending, claims 1-19 are withdrawn as being drawn to

a non-elected invention.

3. Claims 24-26,29-38 are examined on the merits.

4. The text of those sections of Title 35, U.S. Code not included in this action can be

found in a prior Office action.

## **New Rejections**

## Claim Rejections - 35 USC § 112, 1st paragraph

5. Claims 24-26,29-33, and 35-38 are rejected under 35 U.S.C. 112, first paragraph,

as failing to comply with the written description requirement. The claim(s) contains subject

matter which was not described in the specification in such a way as to reasonably convey

to one skilled in the relevant art that the inventor(s), at the time the application was filed, .

had possession of the claimed invention. THIS IS A WRITTEN DESCRIPTION

REJECTION.

The specification teaches that there are numerous peptide sequences which can be

encompassed by the claimed peptide motif of SEQ ID No: 1. Therefore, the sequence

motif of SEQ ID No: 1 encompasses more that represented by sequence such as SEQ ID

No: 9 and 13.

To provide adequate written description and evidence of possession of a claimed genus, the specification must provide sufficient distinguishing identifying characteristics of the genus. The factors to be considered include disclosure of complete or partial structure, physical and/or chemical properties, functional characteristics, structure/function correlation, methods of making the claimed product, or any combination thereof. In this case, the only factor present in the claim is a recitation of

A motif of SEQ ID No: 1, which fails to provide one of skill in the art with a specific core peptide structure. Accordingly, in the absence of sufficient recitation of distinguishing identifying characteristics, the specification does not provide adequate written description of the claimed genus.

Although drawn to DNA arts, the findings in *University of California v. Eli Lilly and Co.*, 119 F.3d 1559, 43 USPQ2d 1398 (Fed. Cir. 1997) and *Enzo Biochem, Inc. v. Gen-Probe Inc.* are relevant to the instant claims. The Federal Circuit addressed the application of the written description requirement to DNA-related inventions in *University Of California v. Eli Lilly and Co.*, 119 F.3d 1559, 43 USPQ2d 1398 (Fed. Cir. 1997). The Court stated that" [a] written description of an invention involving a chemical genus, like a description of a chemical species, 'requires a precise definition, such as by structure, formula, [or] chemical name', of the claimed subject matter sufficient to distinguish it from other materials. "Id. at 1567, 43 USPQ2d at 1405. The court also stated that:

a generic statement such as "vertebrate insulin cDNA" or "mammalian insulin cDNA" without more, is not an adequate written description of the genus because it does not distinguish the genus from others, except by function. It does not specifically define any of the genes that fall within its definition. It does not define any structural features commonly possessed by members of the genus that distinguish them from others. One skilled in the art therefore cannot, as one can do with a fully described genus, visualize or recognize

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the identity of the members of the genus. A definition by function, as we have previously indicated, does not suffice to define the genus because it is only an indication of what the gene does, rather than what it is.

<u>Id.</u> at 1568, 43 USPQ2d at 1406. The court concluded that "naming a type of material generally known to exist, in the absence of knowledge as to what that material consists of, is not a description of that material." <u>Id.</u>

Finally, the court addressed the manner by which a genus of cDNAs might be described. "A description of a genus of cDNAs may be achieved by means of a recitation of a representative number of cDNAs, defined by nucleotide sequence, falling within the scope of the genus or of a recitation of structural features common to the members of the genus, which features constitute a substantial portion of the genus." <u>Id.</u>

The Federal Circuit has recently clarified that a DNA molecule can be adequately described without disclosing its complete structure. See *Enzo Biochem, Inc. V. Gen- Probe Inc.*, 296 F.3d 1316, 63 USPQ2d 1609 (Fed. Cir. 2002). The *Enzo* court adopted the standard that "the written description requirement can be met by show[ing] that an invention is complete by disclosure of sufficiently detailed, relevant identifying characteristics ... i.e., complete or partial structure, other physical and/or chemical properties, functional characteristics when coupled with a known or disclosed correlation between function and structure, or some combination of such characteristics." *Id.* at 1324, 63 USPQ2d at 1613 (emphasis omitted, bracketed material in original).

The inventions at issue in *Lilly* and *Enzo* were DNA constructs *per se*, the holdings of those cases are also applicable to claims such as those at issue here. A disclosure that does not adequately describe a product itself logically cannot adequately describe a method of using that product.

Thus the instant specification may provide an adequate written description of the sequence motif of SEQ ID No: 1, per *Lilly*, by structurally describing representative peptides or by describing "structural features common to the members of the genus, which features constitute a substantial portion of the genus." Alternatively, per *Enzo*, the specification can show that the claimed invention is complete "by disclosure of sufficiently detailed, relevant identifying characteristics, functional characteristics when coupled with a known or disclosed correlation between function and structure, or some combination of such characteristics."

In this case, the specification does not provide one of skill in the art with a core structure of amino acids, instead, the specification provides one of skill in the art with amino acid characteristics (i.e. hydrophobic amino acids) and further attempts to characterize the peptide sequences by claiming a genus by a function alone. Therefore, the claimed sequence motif of SEQ ID No: 1 fails to satisfy either the Lilly or Enzo standards. Although the specification discloses a couple of sequences that fail within the motif, these are not representative of the broad class of peptide sequences that may fall within very large and expansive genus of peptides encompassed by the claimed sequence motif of SEQ ID No: 1. This broad recitation fails to satisfy the standard set out in Enzo because the specification provides no functional characteristics coupled to structural features. Further, the specification also fails to describe the broad genus of peptides encompassed the sequence motif claimed by the test set out in Lilly because the specification describes a limited handful of peptides (i.e. SEQ ID No: 9 and 13 for example). Therefore it fails to describe a representative number of species. Thus the specification does not provide an adequate written description of the claimed invention

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that is required to practice the claimed invention.

## Claim Rejections - 35 USC § 102

6. Claims 24-26,33, and 38 are rejected under 35 U.S.C. 102(b) as being anticiapted by Bayer, E (Peptides: Chem. Biochem., Proc. Amer. Peptide Symp., 1st (1970), meeting date 1968, 99-112). Bayer E teaches an isolated dodecapeptide (AFAFAFAFAF) — (see abstract and attached exhibit 1) Bayer does not characterize the dodecapeptide as having the ability to bind to Hsp47 or that it is capable of generating a cytostatic or cytolytic effect on carcinoma cells, the claimed functional limitation would be an inherent property of the referenced peptide. Moreover, the dodecapeptide is nto full length collagen nor a naturally occurring collagen or fragment thereof. Finally, the dodecapeptide also contains at least one F (Phe) as claimed.

Hence, even though the claims are drawn to a peptide that acts by a specific mechanism, the claimed peptide does not appear to distinguish over the prior art teaching of the same or nearly the same product. The mechanism of action does not have a bearing on the patentability of the invention if the invention was already known or obvious. Mere recognition of latent properties in the prior art does not render nonobvious an otherwise known invention. In re Wiseman, 201 USPQ 658 (CCPA 1979). Granting a patent on the discovery of an unknown but inherent function would remove from the public that which is in the public domain by virtue of its inclusion in, or obviousness from, the prior art. In re Baxter Travenol Labs, 21 USPQ2d 1281 (Fed. Cir. 1991). See M.P.E.P. 2145.

Conclusion

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No claim is allowed. Claim 34 is objected to for depending on a rejected claim.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Christopher H. Yaen whose telephone number is 571-272-0838. The examiner can normally be reached on Monday-Friday 9-5.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Larry Helms, Ph.D. can be reached on 571-272-0832. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

Christopher Yaen Art Unit 1643 October 10, 2006

RIMARY EXAMINER

differed from one another in their amino acid and hydroxy acid content and in their number of ester and amide bonds.

ACCESSION NUMBER:

75:6305 CA

TITLE:

Synthesis of valinomycin analogs with modified side chains and different numbers of amide and ester groups

AUTHOR (S):

Fonina, L. A.; Sanasaryan, A. A.; Vinogradova, E. I. Inst. Khim. Prir. Soedin. im. Shemyakina, Moscow, USSR

CORPORATE SOURCE:

Khimiya Prirodnykh Soedinenii (1971), 7(1), 69-81

SOURCE:

CODEN: KPSUAR; ISSN: 0023-1150

DOCUMENT TYPE:

Journal

LANGUAGE:

Russian

L3 ANSWER 376 OF 377 REGISTRY COPYRIGHT 2006 ACS on STN

RN 26251-07-0 REGISTRY

FS PROTEIN SEQUENCE; STEREOSEARCH

SQL

SEQ

1 AFAFAFAFAF AF

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HITS AT: 1-12

\*\*RELATED SEQUENCES AVAILABLE WITH SEQLINK\*\*

#### REFERENCE 1

Merrifield's solid phase method dets. the sequence of synthetic peptides. AB Failure sequences are low, and polypeptides containing 60-80 amino acids can be synthesized. The purification of the end product is important and the dodecapeptides (Leu-Ala)6 and (Ala-Phe)6 were purified so that no failure sequences could be detected. No racemization of the amino acids occurred.

ACCESSION NUMBER:

73:56410 CA

TITLE:

New results in the solid phase method for the

synthesis of peptides

AUTHOR (S):

Bayer, Ernst

CORPORATE SOURCE:

Dep. of Cem., Univ. of Houston, Houston, TX, USA

SOURCE:

Peptides: Chem. Biochem., Proc. Amer. Peptide Symp., 1st (1970), Meeting Date 1968, 99-112. Editor(s): Weinstein, Boris. Marcel Dekker, Inc.: New York, N.

γ

CODEN: 17XJA8

DOCUMENT TYPE:

Conference

LANGUAGE:

English

### REFERENCE 2

Failure sequences occur during solid phase synthesis of polypeptides, but their number is considerably decreased by acetylation of the amino groups which do not react, or by the use of specially prepared resin-coated glass beads.

ACCESSION NUMBER:

72:101090 CA

TITLE:

SOURCE:

Failure sequences in the solid phase synthesis of

polypeptides

AUTHOR (S):

Bayer, Ernst; Eckstein, H.; Haegele, K.; Koenig,

Wilfried A.; Bruening, W.; Hagenmaier, Hanspaul; Parr,

Wolfgang

CORPORATE SOURCE:

Dep. of Chem., Univ. of Houston, Houston, TX, USA Journal of the American Chemical Society (1970),

92(6), 1735-8

CODEN: JACSAT; ISSN: 0002-7863

DOCUMENT TYPE:

Journal

LANGUAGE:

English

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